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AMENDMENTS TO THE CLAIMS

- 1. (Currently Amended) A method of treating or preventing allergic and inflammatory conditions of the skin or airway passages in a human in need thereof of such treating or preventing while avoiding a food effect associated with other non-sedating antihistamines which comprises orally administering to said human under feed or fasted conditions an amount of desloratedine effective for such treating or preventing while avoiding a food effect on the bioavailability of desloratedine.
- 2. (Original) The method of claim 1 wherein the amount of desloratadine is about 2.5 mg/day to about 45 mg/day.
- 3. (Original) The method of claim 1 wherein the amount of desloratadine is about 2.5 mg/day.
- 4. (Original) The method of claim 1 wherein the amount of desloratadine is about 5 mg/day to about 10 mg/day.
- 5. (Original) The method of claim 1 wherein the amount of desloratadine is about 5 mg/day.
- 6. (Original) The method of claim 1 wherein the desloratadine is administered in a tablet formulation.
- 7. (Original) The method of claim 1 wherein the desloratadine is administered in a syrup formulation.

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8. (Currently Amended) The method of claim 1 wherein the allergic reaction is season seasonal allergic rhinitis, pernninal perennial allergic rhinitis, atopic dermatitis, urticaria or allergic asthma.

- 9. (Currently Amended) A method of treating or preventing allergic and inflammatory conditions of the skin or airway passages in a human in need thereof of such treating or preventing a which comprises orally administering to said human an amount of desloratedine effective for such treating, treating or preventing, while obtaining sustantially substantially the same bioavailability of desloratedine under feed fed or fasted conditions.
- 10. (Original) The method of claim 9 wherein the amount of desloratadine is about 2.5 mg/day to about 45 mg/day.
- 11. (Original) The method of claim 9 wherein the amount of desloratadine is about 2.5 mg/day.
- 12. (Original) The method of claim 9 wherein the amount of desloratadine is about 5 mg/day to about 10 mg/day.
- 13. (Original) The method of claim 9 wherein the amount of desloratadine is about 5 mg/day.
- 14. (Original) The method of claim 9 wherein the desloratadine is administered in a tablet formulation.
- 15. (Original) The method of claim 9 wherein the desloratadine is administered in a syrup formulation.

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16. (Currently Amended) The method of claim 9 wherein the allergic reaction is season seasonal allergic rhinitis, pernninal perennial allergic rhinitis, atopic dermatitis, urticaria or allergic asthma.

- 17. (Currently Amended) A method of treating or preventing seasonal or perennial allergic rhinitis in a human in need thereof of such treating or preventing while avoiding a food effect associated with non-sedating antihistamines which comprises orally administering to said human under feed fed or fasted conditions an amount of desloratedine effective for such treating or preventing while avoiding a food effect on the bioavailability of desloratedine.
- 18. (Original) The method of claim 17 wherein the amount of desloratadine is in the range of about 2.5 mg/day to about 45 mg/day.
- 19. (Original) The method of claim 17 wherein the amount of desloratadine is about 5 mg/day to about 15 mg/day.
- 20. (Original) The method of claim 17 wherein the amount of desloratadine is about 2.5 mg/day.
- 21. (Original) The method of claim 17 wherein the amount of desloratadine is about 5 mg/day.
- 22. (Original) The method of claim 17 wherein the patient is suffering from seasonal allergic rhinitis.
- 23. (Original) The method of claim 17 wherein the patient is suffering from perennial allergic rhinitis.

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24. (Original) The method of claim 17 wherein the desloratadine is administered in a tablet formulation.

- 25. (Original) The method of claim 17 wherein the desloratadine is administered in a syrup formulation.
- 26. (Currently Amended) A method of treating or preventing seasonal or perennial allergic rhinitis in a human in need thereof of such treating or preventing a which comprises orally administering to said human an amount of desloratedine effective for such treating, treating or preventing, while obtaining sustantially substantially the same bioavailability of desloratedine under feed fed or fasted conditions.
- 27. (Original) The method of claim 26 wherein the amount of desloratadine is in the range of about 2.5 mg/day to about 45 mg/day.
- 28. (Original) The method of claim 26 wherein the amount of desloratadine is about 5 mg/day to about 15 mg/day.
- 29. (Original) The method of claim 26 wherein the amount of desloratadine is about 2.5 mg/day.
- 30. (Original) The method of claim 26 wherein the amount of desloratadine is about 5 mg/day.
- 31. (Original) The method of claim 26 wherein the patient is suffering from seasonal allergic rhinitis.

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32. (Original) The method of claim 26 wherein the patient is suffering from perennial allergic rhinitis.

- 33. (Original) The method of claim 26 wherein the desloratadine is administered in a tablet formulation.
- 34. (Original) The method of claim 26 wherein the desloratadine is administered in a syrup formulation.
- 35. (Currently Amended) A method of treating or preventing atopic dermatitis or urticaria in a human in need thereof of such treating or preventing while avoiding a food effect associated with non-sedating antihistamines which comprises orally administering to said human under feed fed or fasted conditions an amount of desloratedine effective for such treating or preventing while avoiding a food effect on the bioavailability of desloratedine.
- 36. (Original) The method of claim 35 wherein the amount of desloratadine is about 2.5 mg/day.
- 37. (Original) The method of claim 35 wherein the amount of desloratadine is about 5 mg/day to about 15 mg/day.
- 38. (Original) The method of claim 35 wherein the amount of desloratadine is about 5 mg/day to about 10 mg/day.
- 39. (Original) The method of claim 35 wherein the amount of desloratadine is about 5 mg/day.

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40. (Original) The method of claim 35 wherein the patient is suffering from atopic dermatitis.

- 41. (Original) The method of claim 35 wherein the patient is suffering from urticaria.
- 42. (Currently Amended) A method of treating or preventing atopic dermatitis or urticaria in a human in need thereof of such treating or preventing a which comprises orally administering to said human an amount of desloratedine effective for such treating, treating or preventing, while obtaining sustantially substantially the same bioavailability of desloratedine under feed fed or fasted conditions.
- 43. (Original) The method of claim 42 wherein the amount of desloratadine is in the range of about 2.5 mg/day to about 45 mg/day.
- 44. (Original) The method of claim 42 wherein the amount of desloratadine is about 2.5 mg/day to about 45 mg/day.
- 45. (Original) The method of claim 42 wherein the amount of desloratadine is about 2.5 mg/day.
- 46. (Original) The method of claim 42 wherein the amount of desloratadine is about 5 mg/day to about 10 mg/day.
- 47. (Original) The method of claim 42 wherein the amount of desloratadine is about 5 mg/day.
- 48. (Original) The method of claim 42 wherein the desloratadine is administered in a tablet formulation.

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49. (Original) The method of claim 42 wherein the desloratadine is administered in a syrup formulation.

- 50. (Original) The method of claim 42 wherein the patient is suffering from atopic dermatitis.
- 51. (Original) The method of claim 42 wherein the patient is suffering from urticaria.